AMENDMENTS TO THE CLAIMS

This listing of claims replaces all prior versions and listings of claims in the application.

Listing of Claims

1. (Currently amended) A compound of formula I

or a pharmaceutically acceptable salt, hydrate, solvate or prodrug of the compound, wherein:

R¹ is hydrogen, -OH, -NO₂, -CN, -COOR, -OCH₂OR, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₁-C₆ alkoxy or halo;

R is C_1 - C_6 alkyl;

R² is a non-radioactive halo or a radioactive halo;

R³ is hydrogen, C₁-C₆ alkyl, C₂-C₆ alkenyl or C₂-C₆ alkynyl; and

R⁴ is hydrogen, C₁-C₆ alkyl, C₂-C₆ alkenyl or C₂-C₆ alkynyl, wherein the alkyl, alkenyl or alkynyl comprises a radioactive carbon or is substituted with a radioactive halo when R² is a non-radioactive halo.

2. (Previously presented) The compound of claim 1, wherein:

R¹ is hydrogen, -OH, -CN, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₁-C₆ alkoxy or halo; and

 R^4 is C_1 - C_6 alkyl, C_2 - C_6 alkenyl or C_2 - C_6 alkynyl, wherein the alkyl, alkenyl or alkynyl comprises a radioactive carbon.

3. (Previously presented) The compound of claim 2, wherein:

R¹ is hydrogen, -OH, -CN, -OCH₃, -CH₃ or -Br; and

R³ is hydrogen or -CH₃; and

$$R^4$$
 is $-^{11}CH_3$.

4. (Currently amended) The compound of claim 1, wherein:

R² is a non-radioactive halo or a radioactive halo, wherein the halo is iodo; and

R⁴ is **hydrogen**, C₁-C₆ alkyl, C₂-C₆ alkenyl or C₂-C₆ alkynyl, wherein the alkyl, alkenyl or alkynyl comprises a radioactive carbon when R² is a non-radioactive halo.

5. (Previously presented) The compound of claim 4, wherein:

the radioactive carbon in R⁴ is ¹¹C.

6. (Currently amended) The compound of claim 5, wherein:

$$R^1$$
 is –OH or C_1 - C_6 alkoxy;

R³ and R⁴ are independently is hydrogen or C₁-C₆ alkyl; and

$$\mathbb{R}^4$$
 is \mathbb{C}_{1} - \mathbb{C}_{6} alkyl.

7. (Currently amended) The compound of claim 6, wherein:

$$R^1$$
 is $-OH$;

$$R^2$$
 is $-^{123}I$ or $-^{125}I$: and

R³ and R⁴ are each is hydrogen.

- 8. (Original) The compound of claim 1, wherein R^2 is a radiofluoro.
- 9. (Currently amended) The compound of claim 8, wherein:

$$R^1$$
 is -OH or C_1 - C_6 alkoxy;

R³ and R⁴-are independently is hydrogen or C₁-C₆ alkyl; and

\mathbb{R}^4 is \mathbb{C}_{1} - \mathbb{C}_{6} alkyl.

10. (Previously presented) The compound of claim 9, wherein:

 R^1 is -OH;

R³ is hydrogen; and

R⁴ is -CH₃.

- 11. (Previously presented) The compound of claim 1, wherein R^4 is C_1 - C_6 alkyl, C_2 - C_6 alkenyl or C_2 - C_6 alkynyl, wherein the alkyl, alkenyl or alkynyl is substituted with a radioactive halo.
- 12. (Previously presented) The compound of claim 11, wherein:

R¹ is -OH or C₁-C₆ alkoxy;

R² is hydrogen;

R³ is hydrogen or C₁-C₆ alkyl; and

 R^4 is C_1 - C_6 alkyl substituted with ^{18}F .

13. (Previously presented) The compound of claim 12, wherein:

 R^1 is -OH;

R³ is hydrogen; and

 R^4 is $-CH_2CH_2CH_2^{18}F$.

- 14. (Original) A pharmaceutical composition comprising
 - (i) an effective amount of a compound of claim 1; and
 - (ii) a pharmaceutically acceptable carrier.

- 15. (Previously presented) A method for detecting amyloid deposit(s) in vivo, comprising:
 - (i) administering to a mammal an effective amount of a compound of claim 1, wherein the compound would bind any amyloid deposit(s) in the mammal; and
 - (ii) detecting binding of the compound to amyloid deposit(s) in the mammal.
- 16. (Previously presented) The method of claim 15, wherein the amyloid deposit(s) is/are located in the brain of the mammal.
- 17. (Previously presented) The method of claim 15, wherein the mammal is a human who is suspected of having Alzheimer's disease, familial Alzheimer's disease, Down's syndrome, Mild Cognitive Impairment or homozygotes for apolipoprotein E4 allele.
- 18. (Previously presented) The method of claim 15, wherein the detecting is effected by gamma imaging, magnetic resonance imaging or magnetic resonance spectroscopy.
- 19. (Previously presented) The method of claim 18, wherein the detecting is effected by gamma imaging.
- 20. (Previously presented) The method of claim 19, wherein the gamma imaging is PET or SPECT.
- 21. (Previously presented) The method of claim 15, wherein the compound is administered intravenously.
- 22. (Previously presented) A method for detecting amyloid deposit(s) in vitro comprising:
 - (i) contacting a bodily tissue with an effective amount of a compound of claim 1, wherein the compound would bind any amyloid deposit(s) in the tissue; and
 - (ii) detecting binding of the compound to amyloid deposit(s) in the tissue.
- 23. (Previously presented) The method of claim 22, wherein the compound is in a solution that further comprises 25-99% ethanol, with the remainder of the solution being water.
- 24. (Previously presented) The method of claim 23, wherein the solution comprises 0-50% ethanol and 0.0001 to 100 μ M of the compound.

- 25. (Previously presented) The method of claim 22 wherein the detecting is effected by bright-field, fluorescence, laser-confocal or cross-polarization microscopy.
- 26. (Previously presented) The method of claim 22, wherein the method further comprises:
 - (iii) separating from the tissue the amyloid deposit(s) bound to the compound; and
 - (iv) quantifying the amyloid deposit(s) bound to the compound.
- 27. (Previously presented) A method for distinguishing an Alzheimer's diseased brain from a normal brain comprising:
 - (i) obtaining tissues from (i) the cerebellum and (ii) another area of the same brain, of a normal mammal and of a mammal suspected of having Alzheimer's disease;
 - (ii) contacting the tissues with a compound of claim 1;
 - (iii) quantifying the amyloid bound to the compound;
 - (iv) calculating the ratio of (a) the amount of amyloid in the area of the brain other than the cerebellum to (b) the amount of amyloid in the cerebellum;
 - (v) comparing the ratio for a normal mammal with the ratio for a mammal suspected of having Alzheimer's disease.